

EXHIBIT C



**UNITED STATES DISTRICT COURT
FOR THE DISTRICT OF MASSACHUSETTS**

IN RE PHARMACEUTICAL INDUSTRY
AVERAGE WHOLESALE PRICE
LITIGATION

MDL No. 1456

CIVIL ACTION: 01-CV-12257-PBS

Judge Patti B. Saris

THIS DOCUMENT RELATES TO
01-CV-12257-PBS AND 01-CV-339

**PLAINTIFFS' AMENDED AND/OR SUPPLEMENTAL REQUEST FOR PRODUCTION
OF DOCUMENTS TO PHASE 1 DEFENDANTS RELATING TO IMS DATA**

Pursuant to Rules 26 and 34 of the Federal Rules of Civil Procedure and the Local Rules of the District Court for the District of Massachusetts, and pursuant to the schedule allowing expert discovery, Plaintiffs hereby request that you produce the documents requested herein within thirty (30) days.

I. DEFINITIONS

1. "Document(s)" is used in the broadest possible sense and means without limitation, any written, printed, typed, photostated, photographed, recorded or otherwise reproduced or stored communication or representation, whether comprised of letters, words, numbers, data, pictures, sounds or symbols, or any combination thereof. This definition includes copies or duplicates of documents contemporaneously or subsequently created which have any non-conforming notes or other markings. Without limiting the generality of the foregoing, "document" includes, but is not limited to, correspondence, memoranda, notes, records, letters,



envelopes, telegrams, messages, studies, analyses, contracts, agreements, working papers, accounts, analytical records, reports and/or summaries of investigations, trade letters, press releases, comparisons, books, calendars, diaries, articles, magazines, newspapers, booklets, brochures, pamphlets, circulars, bulletins, notices, drawings, diagrams, instructions, notes of minutes of meetings or of other communications of any type, including inter-office and intra-office communications, electronic mail/messages and/or “e-mail,” electronically stored telephone messages and/or “voice-mail,” questionnaires, surveys, charts, graphs, photographs, phonograph recordings, films, tapes, disks, data cells, print-outs of information stored or maintained by electronic data processing or word processing equipment, all other data compilations from which information can be obtained (by translation, if necessary, by you through detection devices into usable form), including, without limitation, electromagnetically sensitive storage media such as floppy disks, hard disks and magnetic tapes and any preliminary versions, as well as drafts or revisions of any of the foregoing, whether produced or authored by you or anyone else.

2. “All documents” means every document and every non-identical copy known to you and every such document or writing which you can locate or discover by reasonably diligent efforts, including, but not limited to, documents now in the possession, custody or control of Defendant, its merged or acquired predecessors, its former and present directors, officers, counsel, agents, employees and/or persons acting on its behalf.

3. The term “Defendant” refers to any of the Defendants to whom this is directed, its officers, directors, affiliates, employees, representatives and agents (whether actual, apparent or otherwise).

4. “You” or “Your” means the Defendant responding to these Requests and any of its subsidiaries, divisions, affiliates, officers, directors, employees or agents, including, but not limited to, attorneys and accountants.



II. INSTRUCTIONS

1. A document shall be deemed to be in your control if you have the right to secure the document or copy thereof from another person or public or private entity having possession or custody thereof. If any otherwise responsive document was, but is no longer, in existence or in your possession, custody or control, or has been lost, discarded or destroyed, said document shall be identified as completely as possible including, but not limited to, the following information:

- (a) the date of disposal or disposition from your possession, custody or control;
- (b) the manner of disposal or disposition from your possession, custody or control;
- (c) the reason for disposal or disposition from your possession, custody or control;
- (d) the person authorizing disposal or disposition from your possession, custody or control;
- (e) the document's current or last known custodian;
- (f) the circumstances surrounding the document's disposition from your possession, custody or control;
- (g) the generic category of the document, *e.g.*, memo, letter, computer print-out, etc.;
- (h) the type(s) of information contained in the document; and
- (i) the identity of all persons having knowledge or who had knowledge of the contents of the document.

2. Unless otherwise indicated, the documents to be produced include all documents prepared, sent, dated or received, or those which otherwise came into existence at anytime during the relevant period described herein.



3. (a) Where an objection is made to any document request under Fed. R. Civ. P. 34, the objection shall state with specificity all grounds. Any ground not stated in an objection within the time provided by the Federal Rules of Civil Procedure, or any extensions thereof, shall be waived.

(b) Where a claim of privilege is asserted in objecting to any document demand, or sub-part thereof, and an answer is not provided on the basis of such assertion:

(i) the attorney asserting the privilege shall in the objection to the document demand, or sub-part thereof, identify the nature of the privilege (including work product) that is being claimed and if the privilege is being asserted in connection with a claim or defense governed by state law, indicate the state's privilege rule being invoked; and

(ii) the following information shall be provided in the objection, unless divulgence of such information would cause disclosure of the allegedly privileged information:

(A) for documents: (1) the type of document; (2) general subject matter of the document; (3) the date of the document; and, (4) such other information as is sufficient to identify the document for a subpoena duces tecum, including, where appropriate, the author of the document, the addressee of the document, and, where not apparent, the relationship of the author and addressee to each other;

(B) for oral communications: (1) the name of the person making the communication and the names of persons present while the communication was made and, where not apparent, the relationship of the persons present to the person making the communication; (2) the date and the place of communication; and, (3) the general subject matter of the communication.

4. Notwithstanding the assertion of any objection to production, any document to which an objection is raised containing non-objectional subject matter which is relevant and material to a request must be produced, but that portion of the document for which the objection



is asserted may be withheld or redacted provided that the above-requested information is furnished.

5. This request is continuing and all documents coming into your possession, custody or control which you would have been required to produce had they been available at an earlier time shall be produced forthwith in accordance with the Federal Rules of Civil Procedure.

6. Each document requested herein is requested to be produced in its entirety and without deletion or excisions, regardless of whether you consider the entire document to be relevant or responsive to these requests. If you have redacted any portion of a document, stamp the word "redacted" on each page of the document which you have redacted. Redactions should be included on the privilege log described in Instruction 3.

7. The fact that a document is produced by one defendant does not relieve any other defendant of the obligation to produce his or its copy of the same document, even if the two documents are identical in all respects.

8. In producing documents, you are requested to produce the original of each document requested together with all non-identical copies and drafts of that document. If the original of any document cannot be located, a copy shall be provided in lieu thereof, and shall be legible and bound or stapled in the same manner as the original.

9. All documents shall be produced in the file folder, envelope or other container in which the documents are kept or maintained by you. If, for any reason, the container cannot be produced, produce copies of all labels or other identifying marks.

III. RELEVANT TIME PERIOD

The relevant period of these document requests, unless otherwise indicated, shall be from the launch of the drugs you manufactured that are on Exhibit A, to the date of production and shall include all documents and information which relate in whole or in part to such period, or to events or circumstances during such period, even though dated, prepared, generated or received prior or subsequent to that period.



IV. REQUESTS FOR PRODUCTION

REQUEST FOR PRODUCTION NO. 1:

With respect to the drugs on Exhibit A, the following IMS reports or data for each such drug that you manufactured:

(a) IMS National Disease and Therapeutic Index (NDTI) Data

For each drug that is physician administered and/or covered by Medicare Part B, please provide NDTI data including the following:

- Timeframe: Monthly, from 1991 to the present
- Method of payment
- In terms of TRXs and/or dollars if possible, or in terms of office visits (or “drug uses” or “appearances”) if not
- By manufacturer
- By form (e.g., injectable, tablets, etc.)
- By strength (e.g. 15 mg, 30 mg, etc.)

(b) IMS National Sales Perspective (NSP) (previously the Retail and Provider Perspective)

For each drug that is physician administered and/or covered by Medicare Part B, please provide NSP (or RPP) data including the following:

- Timeframe: Monthly, from 1991 to the present
- Data elements: Units, Extended Units, Dollars
- By channel (e.g. clinic, non-federal hospital, chain, etc.)
- By NDC or its equivalent
- By form
- By strength

**EXHIBIT A****Track 1 Summary of Classification of Drugs by Physician Administered and Part B (8/29/05)**

<i>Manufacturer</i>	<i>Drug Name</i>
Astra Zeneca	Diprivan Pulmicort Zoladex
BMS	Amikin Amphotericin B Blenoxane Coumadin Cytosan Etopophos Fungizone Paraplatin Rubex Taxol Tequin IV Vepesid
GSK	Imitrex Kytril Navelbine Zofran Alkeran Zovirax Retrovir Ventolin Zantac Myleran
J&J	Floxin (Injection only) Haldol (Injection only) Levaquin (IV only) Procrit Remicade
Schering Plough and Warrick	Albuterol Integrilin Intron-A Proventil Temodar



DATED: August 30, 2005

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CERTIFICATE OF SERVICE

I hereby certify that I, Steve W. Berman, an attorney, caused a true and correct copy of the foregoing, **PLAINTIFFS' AMENDED REQUEST FOR PRODUCTION OF DOCUMENTS TO PHASE 1 DEFENDANTS RELATING TO IMS DATA** to be delivered to all counsel of record by electronic service pursuant to Paragraph 11 of the Case Management Order No. 2, by sending on August 30, 2005, a copy to Verilaw Technologies for Posting and notification to all parties

By /s/ Steve W. Berman
Steve W. Berman
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**UNITED STATES DISTRICT COURT
FOR THE DISTRICT OF MASSACHUSETTS**

IN RE PHARMACEUTICAL INDUSTRY
AVERAGE WHOLESAL PRICE
LITIGATION

MDL No. 1456

CIVIL ACTION: 01-CV-12257-PBS

Judge Patti B. Saris

THIS DOCUMENT RELATES TO
01-CV-12257-PBS AND 01-CV-339

**PLAINTIFFS' SECOND AMENDED AND/OR SUPPLEMENTAL REQUEST FOR
PRODUCTION OF DOCUMENTS TO PHASE 1 DEFENDANTS RELATING TO IMS
DATA**

Pursuant to Rules 26 and 34 of the Federal Rules of Civil Procedure and the Local Rules of the District Court for the District of Massachusetts, and pursuant to the schedule allowing expert discovery, Plaintiffs hereby request that you produce the documents requested herein within thirty (30) days. This request does not eliminate the prior IMS requests.

I. DEFINITIONS

1. "Document(s)" is used in the broadest possible sense and means without limitation, any written, printed, typed, photostated, photographed, recorded or otherwise reproduced or stored communication or representation, whether comprised of letters, words, numbers, data, pictures, sounds or symbols, or any combination thereof. This definition includes copies or duplicates of documents contemporaneously or subsequently created which have any non-conforming notes or other markings. Without limiting the generality of the foregoing, "document" includes, but is not limited to, correspondence, memoranda, notes, records, letters,



envelopes, telegrams, messages, studies, analyses, contracts, agreements, working papers, accounts, analytical records, reports and/or summaries of investigations, trade letters, press releases, comparisons, books, calendars, diaries, articles, magazines, newspapers, booklets, brochures, pamphlets, circulars, bulletins, notices, drawings, diagrams, instructions, notes of minutes of meetings or of other communications of any type, including inter-office and intra-office communications, electronic mail/messages and/or “e-mail,” electronically stored telephone messages and/or “voice-mail,” questionnaires, surveys, charts, graphs, photographs, phonograph recordings, films, tapes, disks, data cells, print-outs of information stored or maintained by electronic data processing or word processing equipment, all other data compilations from which information can be obtained (by translation, if necessary, by you through detection devices into usable form), including, without limitation, electromagnetically sensitive storage media such as floppy disks, hard disks and magnetic tapes and any preliminary versions, as well as drafts or revisions of any of the foregoing, whether produced or authored by you or anyone else.

2. “All documents” means every document and every non-identical copy known to you and every such document or writing which you can locate or discover by reasonably diligent efforts, including, but not limited to, documents now in the possession, custody or control of Defendant, its merged or acquired predecessors, its former and present directors, officers, counsel, agents, employees and/or persons acting on its behalf.

3. The term “Defendant” refers to any of the Defendants to whom this is directed, its officers, directors, affiliates, employees, representatives and agents (whether actual, apparent or otherwise).

4. “You” or “Your” means the Defendant responding to these Requests and any of its subsidiaries, divisions, affiliates, officers, directors, employees or agents, including, but not limited to, attorneys and accountants.



II. INSTRUCTIONS

1. A document shall be deemed to be in your control if you have the right to secure the document or copy thereof from another person or public or private entity having possession or custody thereof. If any otherwise responsive document was, but is no longer, in existence or in your possession, custody or control, or has been lost, discarded or destroyed, said document shall be identified as completely as possible including, but not limited to, the following information:

- (a) the date of disposal or disposition from your possession, custody or control;
- (b) the manner of disposal or disposition from your possession, custody or control;
- (c) the reason for disposal or disposition from your possession, custody or control;
- (d) the person authorizing disposal or disposition from your possession, custody or control;
- (e) the document's current or last known custodian;
- (f) the circumstances surrounding the document's disposition from your possession, custody or control;
- (g) the generic category of the document, *e.g.*, memo, letter, computer print-out, etc.;
- (h) the type(s) of information contained in the document; and
- (i) the identity of all persons having knowledge or who had knowledge of the contents of the document.

2. Unless otherwise indicated, the documents to be produced include all documents prepared, sent, dated or received, or those which otherwise came into existence at anytime during the relevant period described herein.



3. (a) Where an objection is made to any document request under Fed. R. Civ. P. 34, the objection shall state with specificity all grounds. Any ground not stated in an objection within the time provided by the Federal Rules of Civil Procedure, or any extensions thereof, shall be waived.

(b) Where a claim of privilege is asserted in objecting to any document demand, or sub-part thereof, and an answer is not provided on the basis of such assertion:

(i) the attorney asserting the privilege shall in the objection to the document demand, or sub-part thereof, identify the nature of the privilege (including work product) that is being claimed and if the privilege is being asserted in connection with a claim or defense governed by state law, indicate the state's privilege rule being invoked; and

(ii) the following information shall be provided in the objection, unless divulgence of such information would cause disclosure of the allegedly privileged information:

(A) for documents: (1) the type of document; (2) general subject matter of the document; (3) the date of the document; and, (4) such other information as is sufficient to identify the document for a subpoena duces tecum, including, where appropriate, the author of the document, the addressee of the document, and, where not apparent, the relationship of the author and addressee to each other;

(B) for oral communications: (1) the name of the person making the communication and the names of persons present while the communication was made and, where not apparent, the relationship of the persons present to the person making the communication; (2) the date and the place of communication; and, (3) the general subject matter of the communication.

4. Notwithstanding the assertion of any objection to production, any document to which an objection is raised containing non-objectional subject matter which is relevant and material to a request must be produced, but that portion of the document for which the objection



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7. The fact that a document is produced by one defendant does not relieve any other defendant of the obligation to produce his or its copy of the same document, even if the two documents are identical in all respects.

8. In producing documents, you are requested to produce the original of each document requested together with all non-identical copies and drafts of that document. If the original of any document cannot be located, a copy shall be provided in lieu thereof, and shall be legible and bound or stapled in the same manner as the original.

9. All documents shall be produced in the file folder, envelope or other container in which the documents are kept or maintained by you. If, for any reason, the container cannot be produced, produce copies of all labels or other identifying marks.

III. RELEVANT TIME PERIOD

The relevant period of these document requests, unless otherwise indicated, shall be from the launch of the drugs you manufactured that are on Exhibit A, to the date of production and shall include all documents and information which relate in whole or in part to such period, or to events or circumstances during such period, even though dated, prepared, generated or received prior or subsequent to that period.



IV. REQUESTS FOR PRODUCTION

REQUEST FOR PRODUCTION NO. 1:

With respect to the drugs on Exhibit A, the following IMS reports or data for each such drug that you manufactured:

(a) IMS National Disease and Therapeutic Index (NDTI) Data

For each drug that is physician administered and/or covered by Medicare Part B, please provide NDTI data including the following:

- Timeframe: Monthly, from 1991 to the present
- Method of payment
- In terms of TRXs and/or dollars if possible, or in terms of office visits (or “drug uses” or “appearances”) if not
- By manufacturer
- By form (e.g., injectable, tablets, etc.)
- By strength (e.g. 15 mg, 30 mg, etc.)

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For each drug that is physician administered and/or covered by Medicare Part B, please provide NSP (or RPP) data including the following:

- Timeframe: Monthly, from 1991 to the present
- Data elements: Units, Extended Units, Dollars
- By channel (e.g. clinic, non-federal hospital, chain, etc.)
- By NDC or its equivalent
- By form
- By strength

**EXHIBIT A****Track 1 Summary of Classification of Drugs by Physician Administered and Part B (8/29/05)**

<i>Manufacturer</i>	<i>Drug Name</i>
Astra Zeneca	Diprivan Pulmicort Zoladex
BMS	Amikin Amphotericin B Blenoxane Coumadin Cytosan Etopophos Fungizone Paraplatin Rubex Taxol Tequin IV Vepesid
GSK	Imitrex Kytril Navelbine Zofran Alkeran Zovirax Retrovir Ventolin Zantac Myleran Lanoxin
J&J	Floxin (Injection only) Haldol (Injection only) Levaquin (IV only) Procrit Remicade Risperdal
Schering Plough and Warrick	Albuterol Integrilin Intron-A Proventil Temodar



DATED: August 31, 2005

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CERTIFICATE OF SERVICE

I hereby certify that I, Steve W. Berman, an attorney, caused a true and correct copy of the foregoing, **PLAINTIFFS' SECOND AMENDED AND/OR SUPPLEMENTAL REQUEST FOR PRODUCTION OF DOCUMENTS TO PHASE 1 DEFENDANTS RELATING TO IMS DATA** to be delivered to all counsel of record by electronic service pursuant to Paragraph 11 of the Case Management Order No. 2, by sending on August 31, 2005, a copy to Verilaw Technologies for Posting and notification to all parties

By /s/ Steve W. Berman
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